

This listing of claims will replace all prior versions, and listings of claims in the application:

**Listing of Claims:**

1-7 (canceled)

1                   8 (previously presented): A method for identifying a compound that modulates  
2 cellular proliferation or chemosensitivity, the method comprising the steps of contacting the  
3 compound with a meiotic recombination 11 (MRE11) polypeptide wherein the polypeptide has at  
4 least 95% amino acid sequence identity to SEQ ID NO:2 and has nuclease activity and  
5 determining a functional effect of said compound by measuring nuclease activity of the MRE11  
6 polypeptide, wherein an effect of said compound on the nuclease activity of said MRE11  
7 polypeptide indicates that said compound modulates cellular proliferation or chemosensitivity.

9-13 (canceled)

1                   14 (previously presented): The method of claim 8, wherein the MRE11  
2 polypeptide is expressed in a eukaryotic host cell.

15-22 (canceled)

1                   23 (original): The method of claim 8, wherein modulation is inhibition of  
2 cellular proliferation.

1                   24 (original): The method of claim 8, wherein modulation is inhibition of cancer  
2 cell proliferation.

1                   25 (original): The method of claim 8, wherein modulation is activating  
2 sensitivity to chemotherapeutic reagents.

1                   26 (original): The method of claim 8, wherein modulation is activating  
2 sensitivity of cancer cells to chemotherapeutic reagents.

1                   27 (original): The method of claim 14, wherein the host cell is a cancer cell.

1                   28 (original): The method of claim 27, wherein the cancer cell is a breast,  
2 prostate, colon, or lung cancer cell.

1                   29 (original): The method of claim 27, wherein the cancer cell is a transformed  
2 cell line.

1                   30 (original): The method of claim 29, wherein the transformed cell line is PC3,  
2 HI299, MDA-MB-231, MCF7, A549, or HeLa.

1                   31 (previously presented): The method of claim 27, wherein the cancer cell is a  
2 p53 null or mutant cell.

1                   32 (previously presented): The method of claim 27, wherein the cancer cell is a  
2 p53 wild-type cell.

1                   33 (original): The method of claim 27, wherein the cancer cell is treated with  
2 bleomycin or etoposide.

1                   34 (original): The method of claim 8, wherein the polypeptide is recombinant.

1                   35 (original): The method of claim 8, wherein the polypeptide is encoded by a  
2 nucleic acid having a sequence of SEQ ID NO:1.

1                   36 (original): The method of claim 8, wherein the compound is an antibody.

1                   37 (original): The method of claim 8, wherein the compound is an antisense  
2 molecule.

1                   38 (original): The method of claim 8, wherein the compound is a small organic  
2 molecule.

1                    39 (original): The method of claim 8, wherein the compound is a peptide.

1                    40 (original): The method of claim 39, wherein the peptide is circular.

41-52 (canceled)

1                    53 (previously presented): The method of claim 8, wherein the MRE11  
2 polypeptide has an amino acid sequence of SEQ ID NO:2.

1                    54 (previously presented): The method of claim 8, wherein the MRE11  
2 polypeptide is encoded by a nucleic acid sequence having at least 95% nucleic acid sequence  
3 identity to SEQ ID NO:1.